

Respondents	Number of respondents	Number of responses/re-spondent	Avg. burden/response (in hrs.)	Total burden (in hrs.)
General Public	1200	1	.1666	200

3. X-Ray Examination Program—(0920-0020)—Extension

The X-ray Examination Program is a federally mandated program under the Federal Mine Safety and Health Act of 1977, PL-95-164. The Act provides the regulatory guidance for the

administration of the National Coal Workers' X-ray Surveillance Program, a surveillance program to protect the health and safety of underground coal miners. This program requires the gathering of information from coal mine operators, participating miners, participating x-ray facilities and

participating physicians. The Appalachian Laboratory for Occupational Safety and Health (ALOSH), National Institute for Occupational Safety and Health (NIOSH) is charged with administration of this program. Total annual burden hours are 4,791.

Form title	Total of respondents	Responses/re-spondent	Avg. burden/response (in hrs.)	Total burden (in hrs.)
Roentgenographic Interpretation Title 42 CFR 37.40 (b) 37.50(a); 37.60(a)	20,000	1	0.05	1,000
Miner Identification Title 42 CFR 37.20 37.40(b); 37.60(a)	10,000	1	0.333	3,333
Coal Mine Operator Plan Title 42 CFR 37.4(a)	500	1	0.5	250
Facility Certification Title 42 CFR 37.42(c)	300	1	0.5	150
Interpreting Physician Certification Title 42 CFR 37.51(c)	350	1	0.1666	58

Dated: February 24, 1998.

Kathy Cahill,

Associate Director for Policy Planning and Evaluation, Centers for Disease Control and Prevention (CDC).

[FR Doc. 98-5662 Filed 3-4-98; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Dermatologic and Ophthalmic Drugs Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). At least one portion of the meeting will be closed to the public.

Name of Committee: Dermatologic and Ophthalmic Drugs Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA regulatory issues.

Date and Time: The meeting will be held on March 19 and 20, 1998, 8:30 a.m. to 5:30 p.m.

Location: Holiday Inn, Walker Room, Two Montgomery Village Ave., Gaithersburg, MD.

Contact Person: Tracy Riley or Angie Whitacre, Center for Drug Evaluation and Research (HFD-21), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-5455, or FDA Advisory Committee Information

Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 12534. Please call the Information Line for up-to-date information on this meeting.

Agenda: On March 19, 1998, the committee will discuss generic topical dermatologicals draft guidance. On March 20, 1998, the committee will participate in a scientific discussion of clinical trial design questions for products intended for the treatment of psoriasis.

Procedure: On March 19, 1998, from 8:30 a.m. to 1 p.m., and on March 20, 1998, from 8:30 a.m. to 5:30 p.m., the meeting is open to the public. Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions should be made to the contact person by March 11, 1998. Oral presentations from the public will be scheduled between approximately 8:30 a.m. and 9 a.m., on March 19 and 20, 1998. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before March 13, 1998, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Closed Committee Deliberations: On March 19, 1998, from 1 p.m. to 5:30 p.m., the meeting will be closed to permit discussion and review of trade secret and/or confidential information (5 U.S.C. 552b(c)(4)). This portion of the

meeting will be closed to permit discussion on pending investigational new drug applications issues.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: February 25, 1998.

Michael A. Friedman,

Deputy Commissioner for Operations.

[FR Doc. 98-5630 Filed 3-4-98; 8:45 am]

BILLING CODE 4160-01-F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 95N-0309]

Agency Information Collection Activities; Announcement of OMB Approval

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Infant Formula Requirements" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (the PRA).

FOR FURTHER INFORMATION CONTACT: Margaret R. Schlosburg, Office of Information Resources Management (HFA-250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-1223.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of December 19, 1997